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WHAT IS CLAIMED IS:

1	 A process for producing a magnetic anastomotic component suitable 			
2	for implantation in a patient's body, the process comprising steps of:			
3	forming an anastomotic component having a desired configuration from a			
4	material capable of producing a magnetic field, the anastomotic component having an exterior			
5	surface;			
6	processing the anastomotic component to make the exterior surface suitable			
7	for receiving a layer of biocompatible material; and			
8	providing the exterior surface of the anastomotic component with a layer of			
9	biocompatible material.			
1	2. The process of claim 1 wherein the processing step is performed to			

- make the exterior surface of the component substantially smooth.
- The process of claim 2 wherein the processing step comprises 3. removing unwanted material from the exterior surface of the component by abrasive microblasting.
- The process of claim 3 wherein the processing step comprises placing 4. the component in a mechanically abrasive environment.
- The process of claim 2 wherein the processing step comprises grinding 5. 1 the exterior surface of the component. 2
 - The process of claim 2 wherein the processing step comprises acid 6. etching the exterior surface of the component.
- The process of claim 1 wherein the providing step comprises disposing 7. 1 a layer of biocompatible material over another layer of material that covers the exterior 2 surface of the anastomotic component. 3
- The process of claim 7 wherein the layer of biocompatible material is 1 Gold and the other layer of material is Gold or Nickel. 2
 - 9. The process of claim 1, further comprising electropolishing the component after placing a final layer of material thereon.

1	10.	The process of claim 1 wherein the component has an overall thickness		
2	within the range of fr	om about 0.010 to about 0.030 inch, and the biocompatible layer has a		
3	thickness within the r	range of from about 0.0002 to about 0.0020 inch.		
1	11.	The process of claim 1 wherein the component is formed from NeoFeB		
2	and a layer of biocom	apatible material is placed over the NeoFeB.		
1	12.	The process of claim 1 wherein a portion of the exterior surface is		
2	formed with means for	or enhancing engagement between the component and the tissue of a		
3	vessel.			
1	13.	The process of claim 1 wherein the forming step forms a component		
2	comprised entirely of	a material capable of producing a magnetic field.		
1	14.	The process of claim 1 wherein the forming step forms a component		
2		ration and the processing step changes the component to a second		
3	configuration having structural differences from the first configuration.			
3	configuration having	structural differences from the first configuration.		
1	15.	The process of claim 1 wherein the providing step comprises plating		
2	the exterior surface o	f the component.		
1	16.	The process of claim 15 wherein the exterior surface of the component		
2	is plated more than o	nce.		
1	17.	The process of claim 1 wherein further comprising assembling the		
2	anastomotic compone	ent is assembled with a delivery device for packaging and sterilization.		
1	18.	The process of claim 1 wherein the anastomotic component is		
2	packaged and steriliz	ed after the providing step.		
1	19.	The process of claim 18 wherein the component is magnetized either		
2	before or after being	packaged and sterilized.		
1	20.	A process for producing a magnetic anastomotic component suitable		
2	for implantation in a	patient's body, the process comprising steps of:		

forming an anastomotic component having a desired configuration from a

4	material capable of producing a magnetic field;			
5	packaging the component;			
6	sterilizing the component; and			
7	magnetizing the component in the package.			
1	21.	The process of claim 20 wherein the anastomotic component is		
2	packaged, magnetized and then sterilized.			
1	22.	The process of claim 21 wherein the component is packaged, sterilized		
2	and then magnetized.			
1	23.	The process of claim 22 wherein the component is sterilized by gas.		
1	24.	The process of claim 21 wherein the packaging step comprises		
2	including a plurality of magnetic anastomotic components as part of a kit.			
		C. 1. 'OA - Level's the moderning stan further comprises		
1	25.	The process of claim 24 wherein the packaging step further comprises		
2	including at least one delivery device in the kit.			
1	26.	The process of claim 20 further comprising microblasting or acid-		
2	etching an exterior surface of the component to remove unwanted material, and then coating			
3	the compatible with a layer of biocompatible material prior to the packaging step.			
1	27.	A process for producing a magnetic anastomotic component suitable		
2	for implantation in a	patient's body, the process comprising steps of:		
3	provid	ding an anastomotic component having an ability to produce a magnetic		
4	field, the component	having an exterior surface;		
5	placir	ng a layer of material on a first portion of the exterior surface of the		
6	component so as to leave a second portion of the exterior surface of the component uncovered			
7	by the material; and			
8	magn	etizing the component.		
1	28.	The process of claim 27 wherein the material placed on the first portion		
2	of the exterior is par	amagnetic.		
1	29.	The process of claim 28 wherein the second portion of the exterior		
2	surface of the compo	onent defines an area of concentrated magnetic flux.		

1		30.	The process of claim 29 further comprising placing a layer of different		
2	material over the exterior surface of the component.				
1		31.	The process of claim 30 wherein the different material has diamagnetic		
2	properties.	51.	The process of claims of material and characters material map characters		
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1		32.	The process of claim 29 wherein the second portion of the component		
2	defines a continuous area of concentrated flux.				
1		33.	A process for producing a magnetic anastomotic component suitable		
2	for implantati	on in a	patient's body, the process comprising steps of:		
3		formi	ng an anastomotic component having a desired configuration from a		
4	material capable of producing a magnetic field, the component having an exterior surface;				
5		subjec	cting the component to an acid etching process to remove surface		
6	irregularities;	and			
7	J	provid	ling the exterior surface of the component with a layer of biocompatible		
8	material.	1			
1		34.	The process of claim 22 wherein the subjecting step is performed by		
1	1		The process of claim 33 wherein the subjecting step is performed by		
2	placing the co	ompone	nt in a solution containing phosphoric acid.		
1		35.	The process of claim 34 wherein the component is placed in the		
2	phosphoric ac	cid solu	tion for an amount of time within the range of from about 5 minutes to		
3	about 15 min	utes.			
1		36.	The process of claim 34 further comprising subjecting the solution to		
2	alactric noton		or the soid etching sten		

- electric potential after the acid etching step
- 37. The process of claim 33 further comprising providing at least a portion of the exterior surface of the component with traction structure for enhancing engagement between the component and the tissue of a vessel.
- 1 38. The process of claim 37 wherein the traction structure comprises a surface of the component provided with adhesive.

- 1 39. The process of claim 37 wherein the traction structure comprises a 2 surface of the component provided with tissue-gripping elements configured to grip the tissue 3 of a vessel.
- 1 40. The process of claim 37 wherein the traction structure comprises a 2 surface of the component provided with a tacky coating configured to stick to vessel tissue.